

	Type	L #	Hits	Search Text	DBs	Time Stamp	Comments	Error Definition	Errors
1	BRS	L1	290	echinocandin	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:32			0
2	BRS	L3	28828	antifungal	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:32			0
3	BRS	L4	13	2 same 3	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:33			0
4	BRS	L5	5159	anti-fungal	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:33			0
5	BRS	L6	4	2 same 5	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:37			0
6	BRS	L7	1	wo-9613272-\$.did.	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 09:37			0
7	BRS	L2	24	echinocandin adj derivative	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:07			0
8	BRS	L8	166	1 same (3 or 5)	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:36			0
9	BRS	L9	36	fauveau adj patrick.in.	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:37			0
10	BRS	L10	4	hawser adj stephen.in.	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:38			0
11	BRS	L11	0	lebourg adj gilles.in.	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:39			0
12	BRS	L12	9	schio adj laurent.in.	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:39			0
13	BRS	L13	4	(9 or 10 or 12) and 2	USPAT; US-PGPUB; EPO; JPO; DERWENT	2003/09/03 10:40			0

=> d his

(FILE 'HOME' ENTERED AT 15:19:38 ON 03 SEP 2003)

FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH, AGRICOLA'  
ENTERED AT

15:20:04 ON 03 SEP 2003

L1 1691 S ECHINOCANDIN  
L2 379 S L1 (P) DERIVATIVE  
L3 115 S ECHINOCANDIN DERIVATIVE  
L4 121417 S ANTI-FUNGAL OR ANTIFUNGAL  
L5 159 S (L2 OR L3 ) (P) L4  
L6 72 DUPLICATE REMOVE L5 (87 DUPLICATES REMOVED)  
L7 18 S FAUVEAU PATRICK/AU  
L8 15 S HAWSER STEPHEN/AU  
L9 1 S LEBOURG GILLES/AU  
L10 22 S SCHIO LAURENT/AU  
L11 5 S (L10 OR L7 OR L8 OR L9) AND L6

The STN search turned up 36 cpds in 2 citations  
 only 18 cpds were in the 2 cites. these are the

KAM 10/018,073

=> d 145 1-18

L45 ANSWER 1 OF 18 REGISTRY COPYRIGHT 2003 ACS on STN

RN 310462-00-1 REGISTRY

CN Deoxymulundocandin, 1-[(4S)-4-[[[(1R,2R)-2-aminocyclohexyl]amino]-N2-[[4''-(pentyloxy)[1,1':4',1''-terphenyl]-4-yl]carbonyl]-L-ornithine]- (9CI) (CA INDEX NAME)

FS PROTEIN SEQUENCE; STEREOSEARCH

MF C63 H83 N9 O14

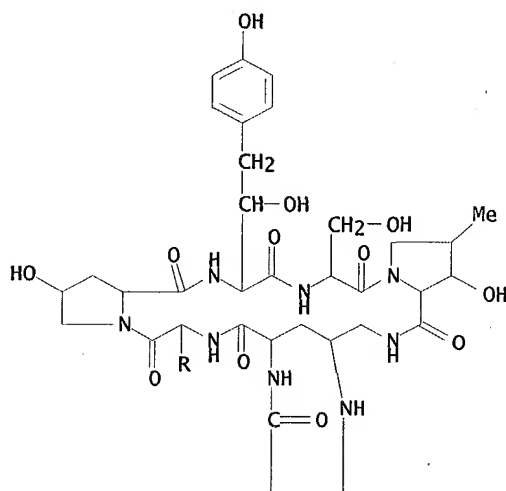
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SR CA

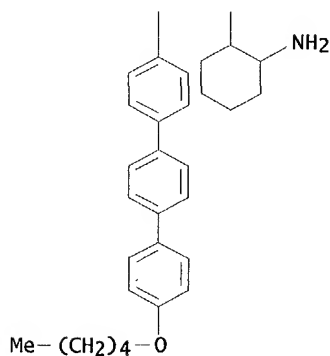
\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

Registry records for the  
 remaining 18. The Reg # are  
 really high - so  
 these must be  
 pretty new & not  
 yet assigned to  
 a citation.

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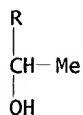


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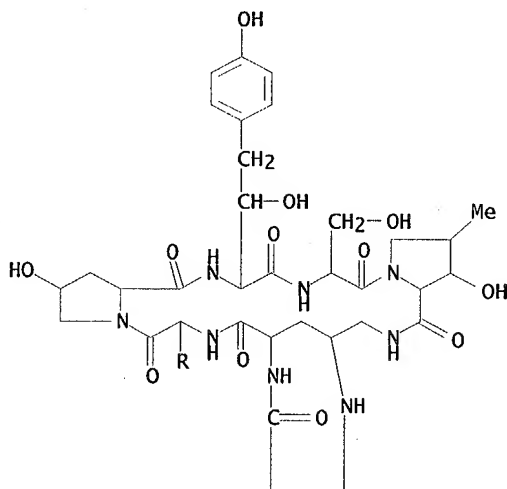
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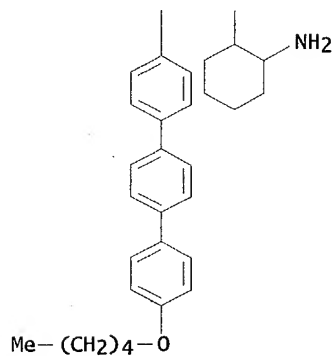
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RN 310461-98-4 REGISTRY  
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FS PROTEIN SEQUENCE; STEREOSEARCH  
MF C63 H83 N9 O14  
CI COM  
SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

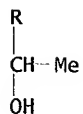
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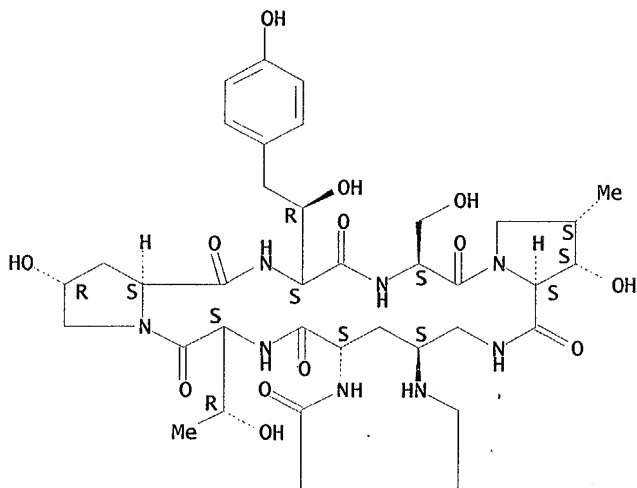


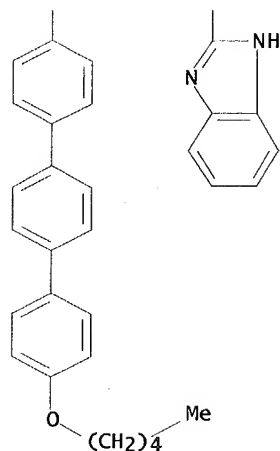
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 RN 310461-96-2 REGISTRY  
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C65 H78 N10 O14  
 CI COM  
 SR CA

\*\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

Absolute stereochemistry.

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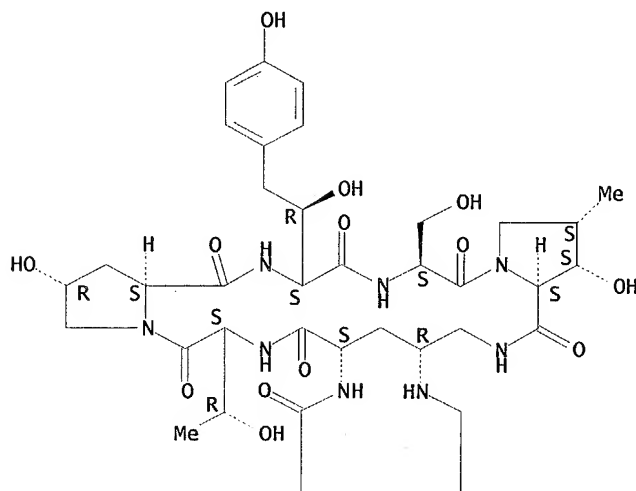


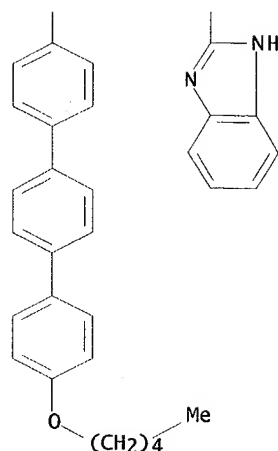


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 MF C65 H78 N10 O14  
 CI COM  
 SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

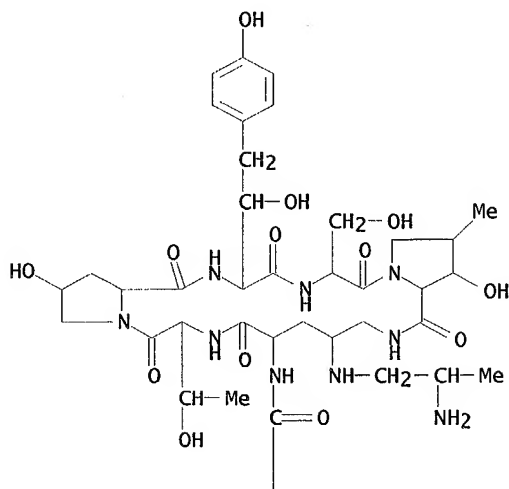
Absolute stereochemistry.



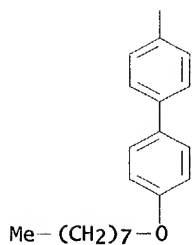


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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C57 H81 N9 O14  
 CI COM  
 SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*



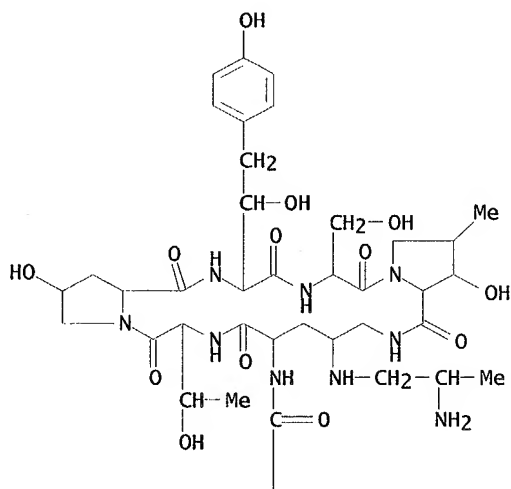
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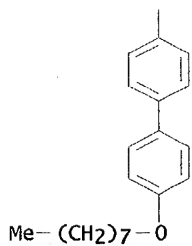
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

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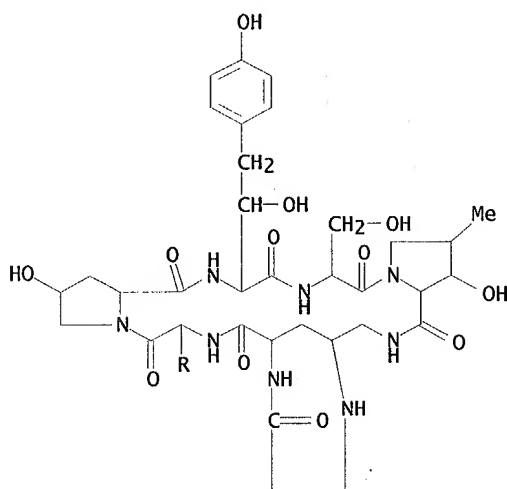


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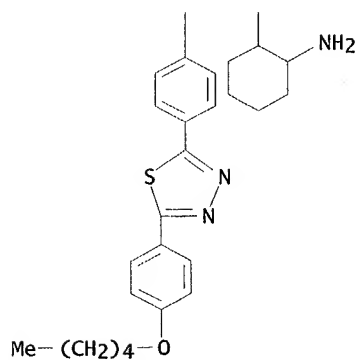
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RN 310459-69-9 REGISTRY  
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MF C59 H79 N11 O14 S  
CI COM  
SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

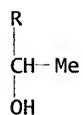
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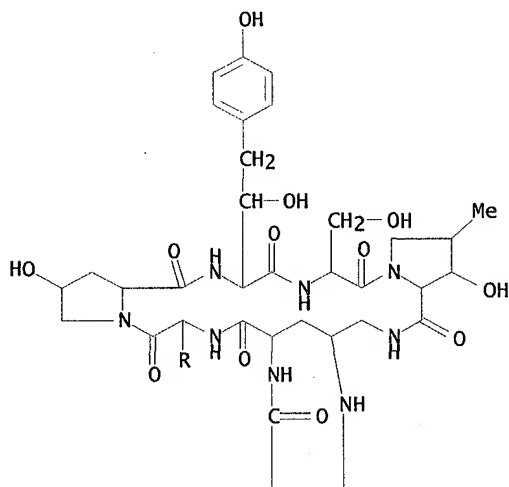
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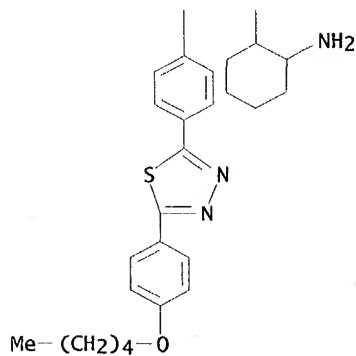
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 CI COM  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

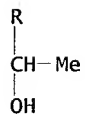
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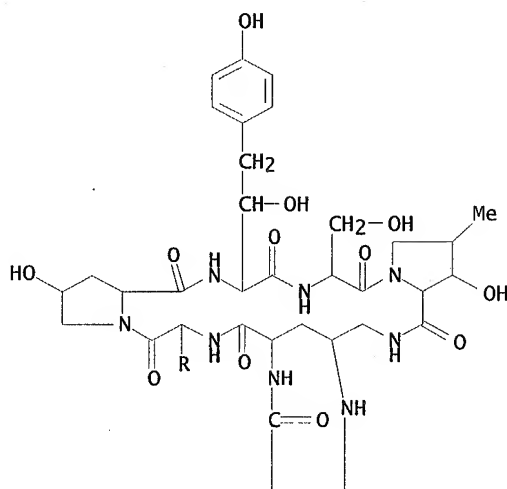
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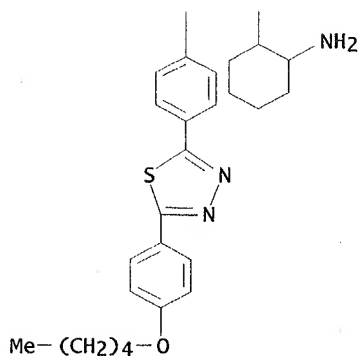
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 (CA INDEX NAME)  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

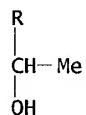
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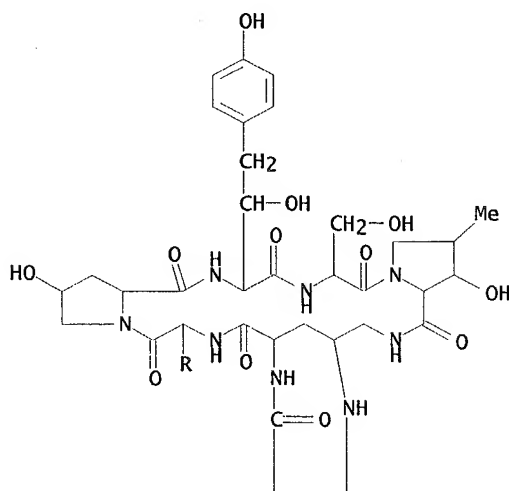
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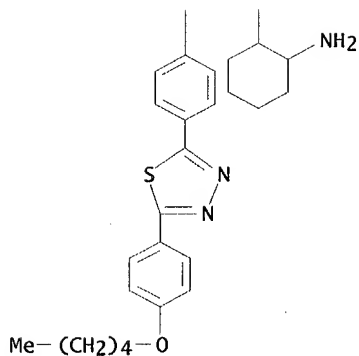
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 RN 310459-57-5 REGISTRY  
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 (CA INDEX NAME)  
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 CI COM  
 SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

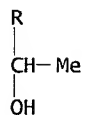
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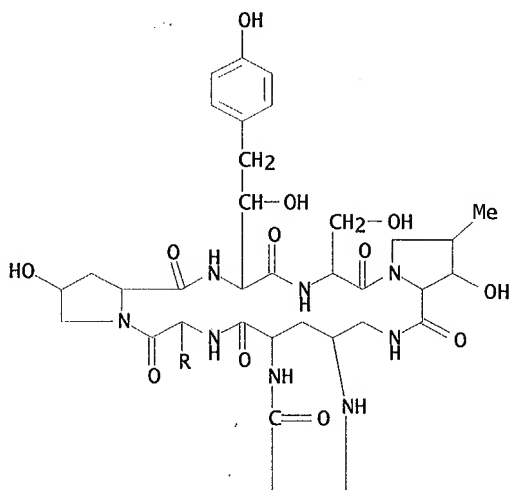
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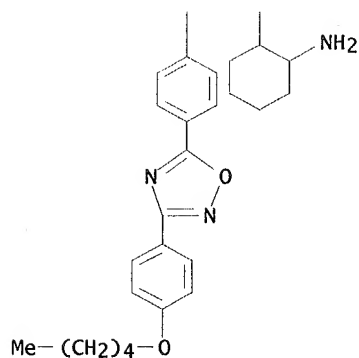
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 RN 310459-41-7 REGISTRY  
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 (CA INDEX NAME)  
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 MF C59 H79 N11 O15  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

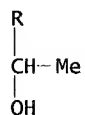
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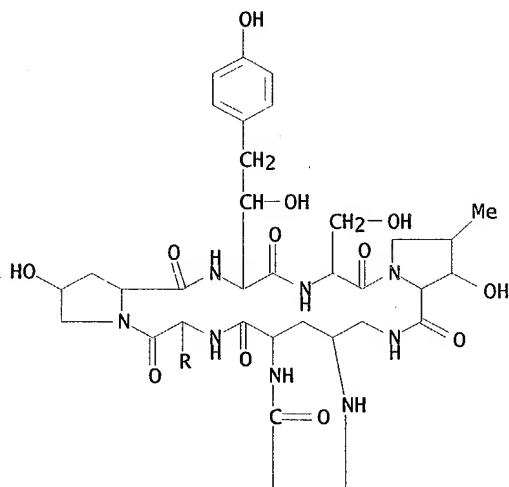
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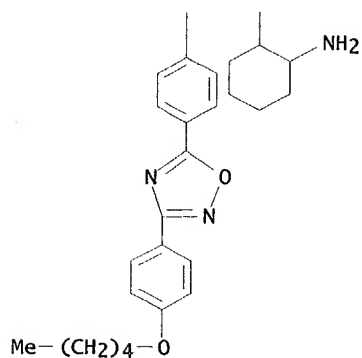
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\*\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

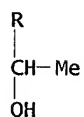
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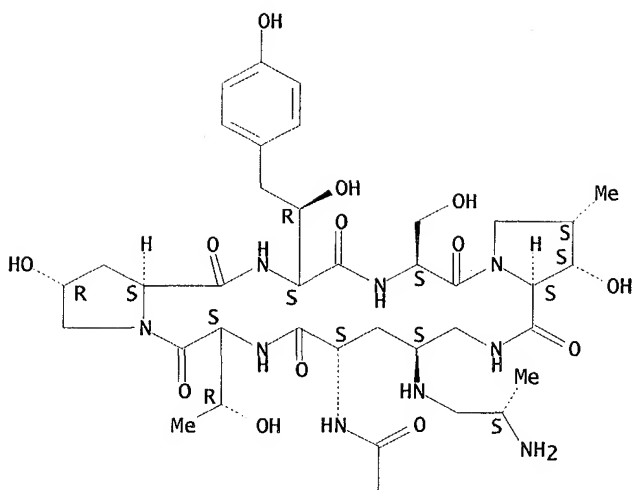


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 RN 310459-35-9 REGISTRY  
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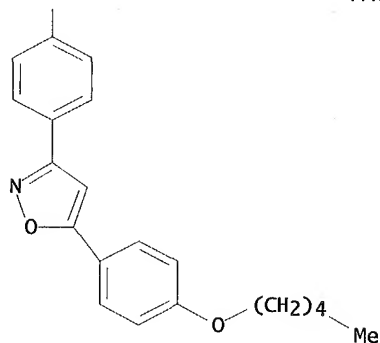
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Absolute stereochemistry.

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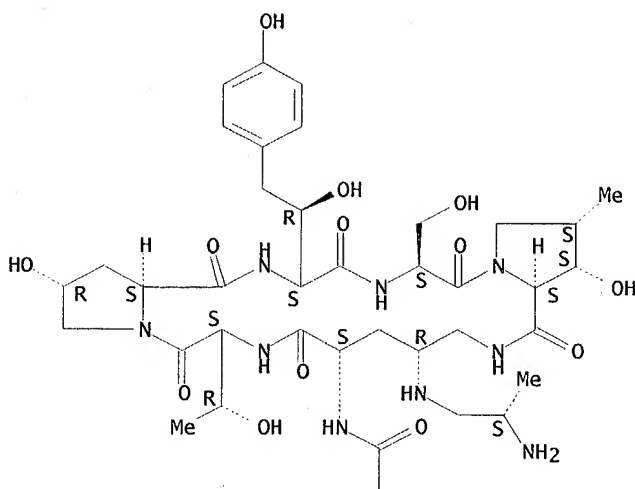
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C57 H76 N10 O15  
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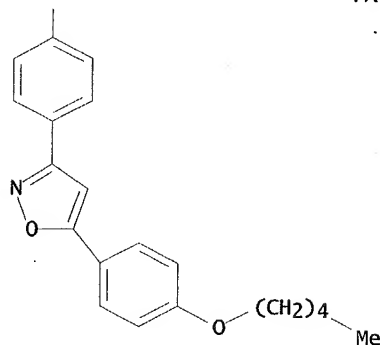
Absolute stereochemistry.



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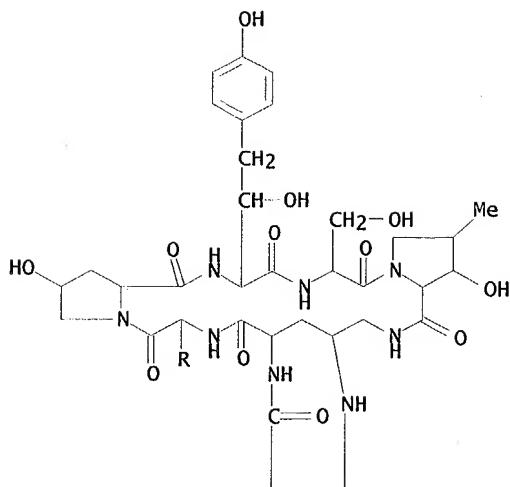
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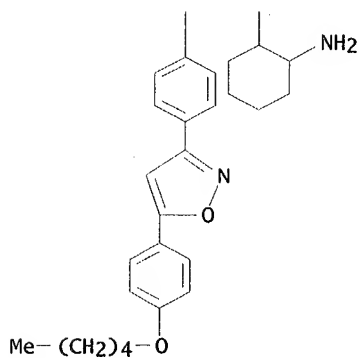
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C60 H80 N10 O15  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

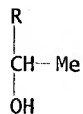
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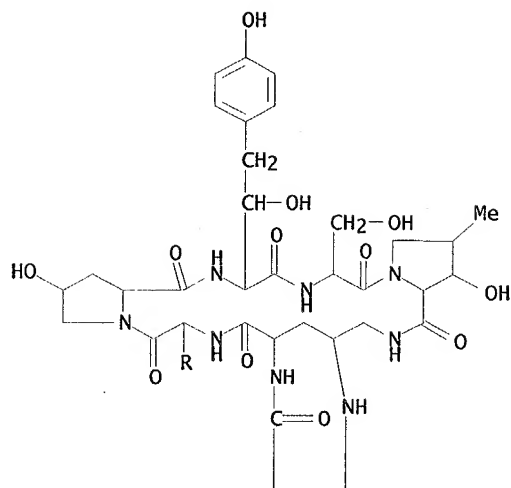
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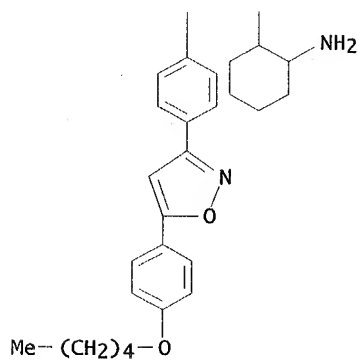
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 RN 310459-26-8 REGISTRY  
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C60 H80 N10 O15  
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\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

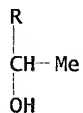
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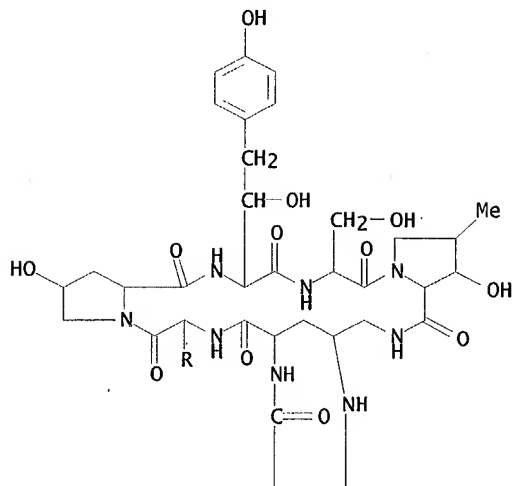
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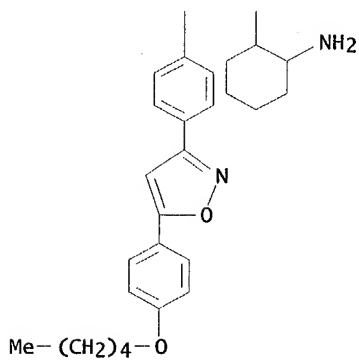
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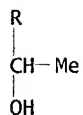
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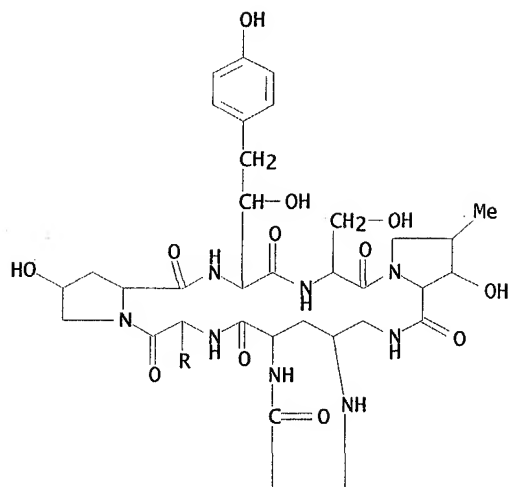
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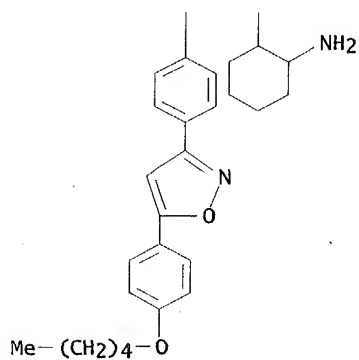
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 FS PROTEIN SEQUENCE; STEREOSEARCH  
 MF C60 H80 N10 O15  
 CI COM  
 SR CA

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

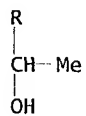
PAGE 1-A



PAGE 2-A



PAGE 3-A



# Inventu Search

KAM 10/018,073

=> d que 110

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L9 12 SEA FILE=REGISTRY ABB=ON PLU=ON L8 AND "DEOXYMULUNDOCANDIN" ← 12 cpds w/ claimed  
L10 1 SEA FILE=CAPLUS ABB=ON PLU=ON L7 AND L9 1 cite ring str

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L10 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2000:881186 CAPLUS

DOCUMENT NUMBER: 134:17731

TITLE: Echinocandin derivatives, method for preparing same and application as glucan synthase inhibitors and antifungal agents

INVENTOR(S): Fauveau, Patrick; Hawser, Stephen; Lebourg, Gilles; Schio, Laurent

PATENT ASSIGNEE(S): Hoechst Marion Roussel, Fr.

SOURCE: PCT Int. Appl., 24 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: French

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PCT document  
Parent case

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
FR 2794746	A1	20001215	FR 1999-7251	19990609
FR 2794746	B1	20021206		
EP 1189933	A1	20020327	EP 2000-942169	20000608
EP 1189933	B1	20030409		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2003501441	T2	20030114	JP 2001-502458	20000608
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PRIORITY APPLN. INFO.: FR 1999-7251 A 19990609				
WO 2000-FR1568 W 20000608				
OTHER SOURCE(S): CASREACT 134:17731; MARPAT 134:17731				
GI				

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB The invention concerns in all possible isomeric forms as well as their mixts., cyclic peptides I wherein: R represents a linear, branched or cyclic chain; either R1 represents H or CH3 and R2 represents cyclohexyl substituted by an amine, cyanoalkyl; or R1 and R2 form with the nitrogen which bears them a cycle with 3, 4 or 5 carbons optionally substituted by an amine; R3 represents hydrogen, Me or hydroxyl; R4 represents hydrogen or hydroxyl; T represents hydrogen, Me, CH2CONH2, CH2CN, a (CH2)2NH2 or (CH2)2Nalk+X- radical, X being halogen and alk an alkyl radical; Y

KAM 10/018,073

represents hydrogen, hydroxyl, halogen or OSO<sub>3</sub>H; W represents H or OH; Z represents H, CH<sub>3</sub>. The compds. of formula I have antifungal properties. Thus, Trans 1-[4-[(2-aminocyclohexyl)amino]-N<sub>2</sub>-[[4'-(pentyloxy)[1,1':4',1''terphenyl]-4-yl]carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate was prepd. and tested for its inhibition of glucan synthase of *Candida albicans* and of the enzyme prepd. from *Aspergillus fumigatus*.

IT 310461-86-0P 310461-89-3P 310461-95-1P

310461-97-3P 310461-99-5P 310462-01-2P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(echinocandin derivs., method for prep. same and application as glucan synthase inhibitors and antifungal agents)

RN 310461-86-0 CAPLUS

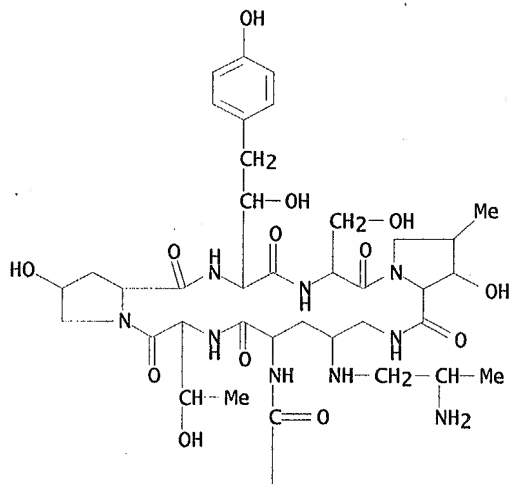
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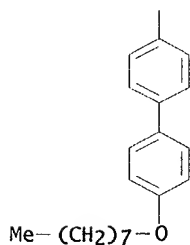
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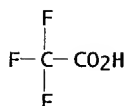
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KAM 10/018,073

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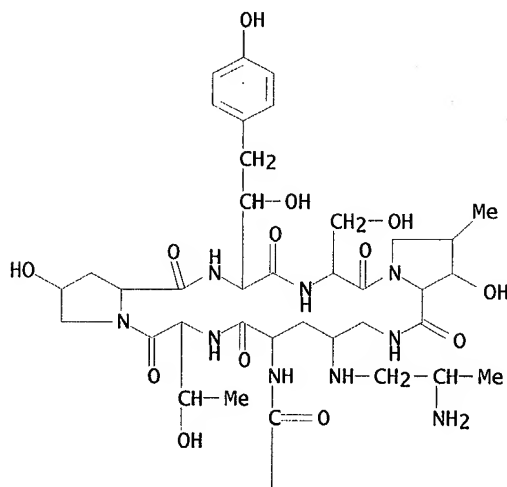


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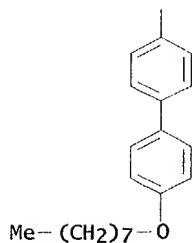
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PAGE 1-A



PAGE 2-A



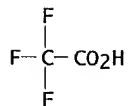
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KAM 10/018,073

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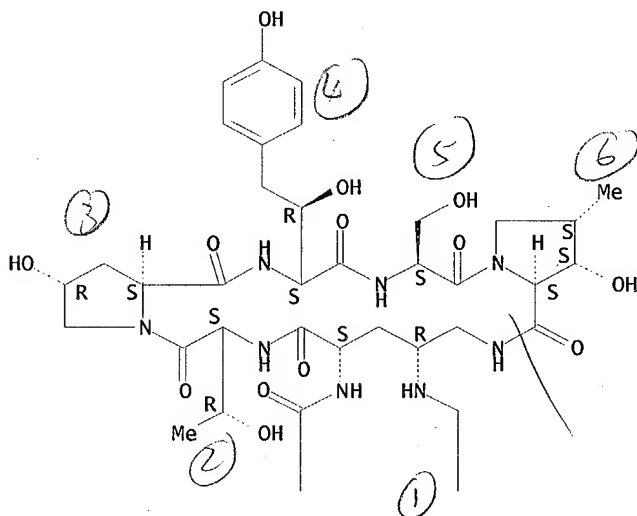
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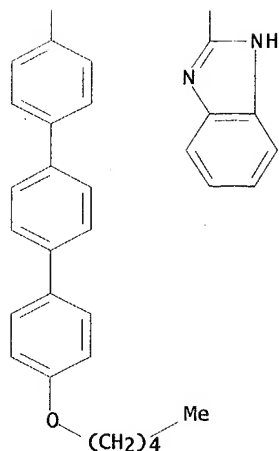
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Absolute stereochemistry.

PAGE 1-A

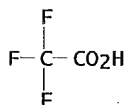


*this could be #1 in  
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or  
next page*



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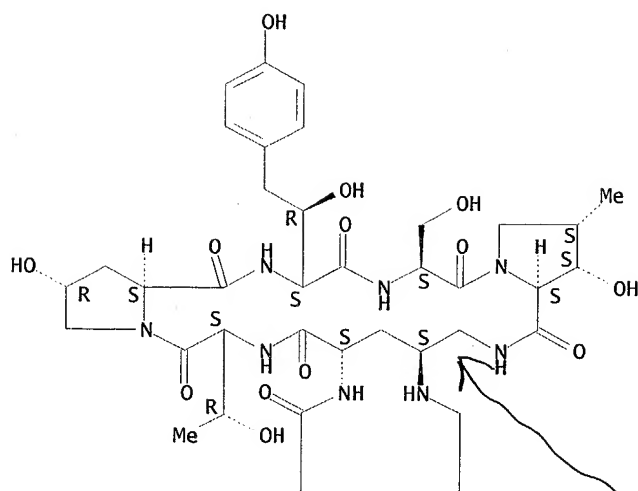
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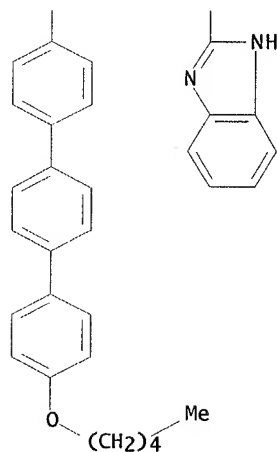
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Absolute stereochemistry.

PAGE 1-A



PAGE 2-A

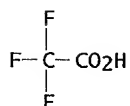


this could be #1 in  
claim 15

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CM 2

CRN 76-05-1  
CMF C2 H F3 O2



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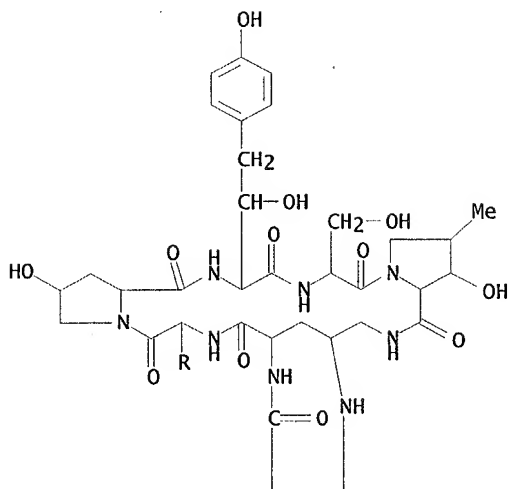
KAM 10/018,073

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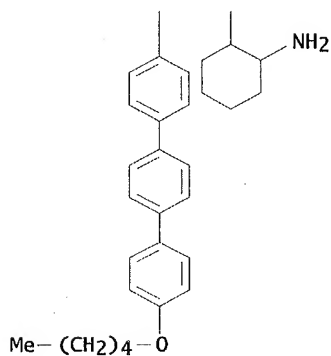
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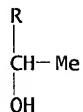
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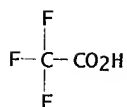


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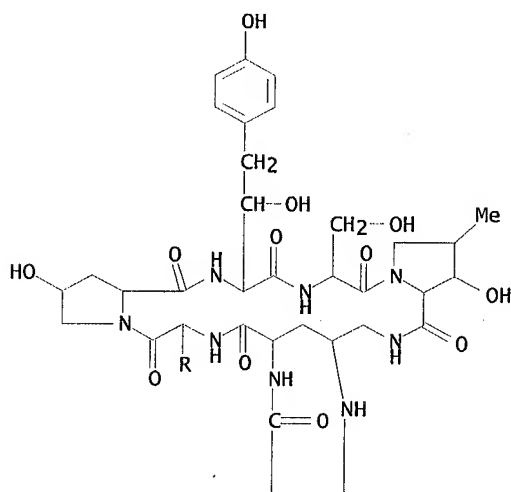


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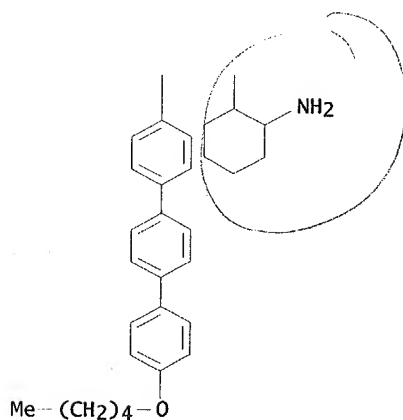
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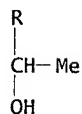
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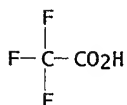
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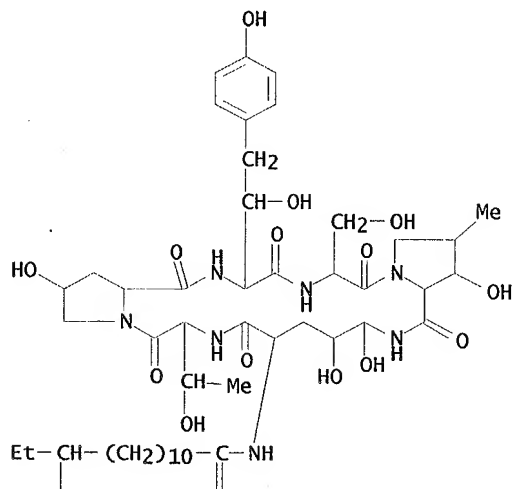


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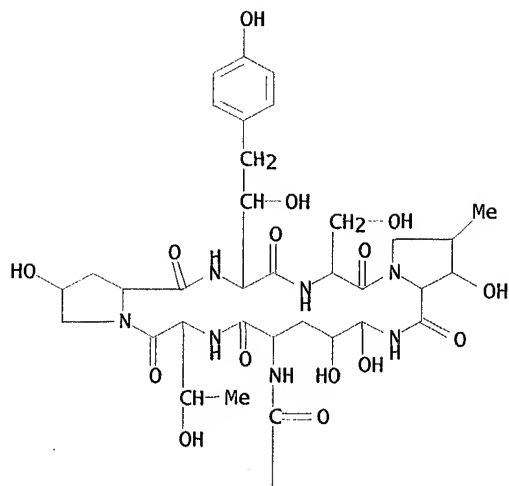


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 RN 138626-63-8 CAPLUS  
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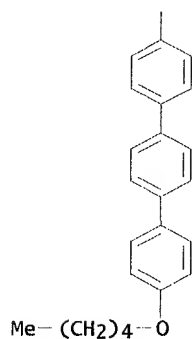


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PAGE 1-A



PAGE 2-A



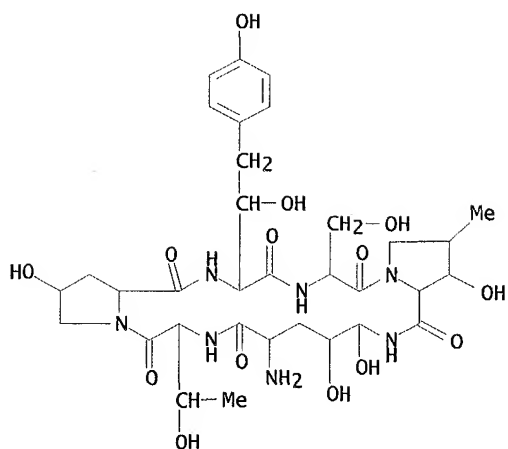
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(echinocandin derivs., method for prepg. same and application as glucan  
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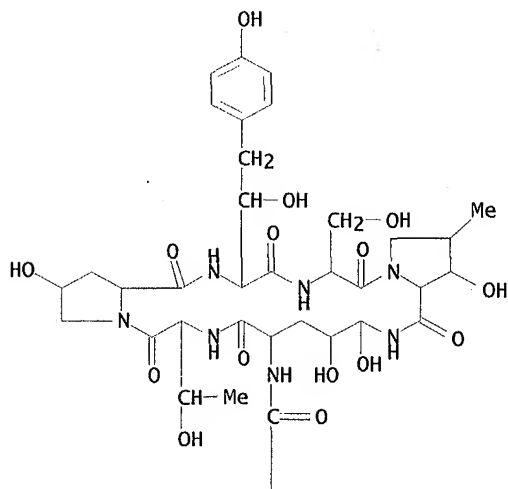
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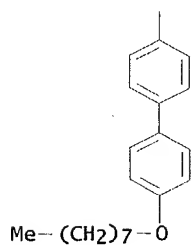


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PAGE 1-A



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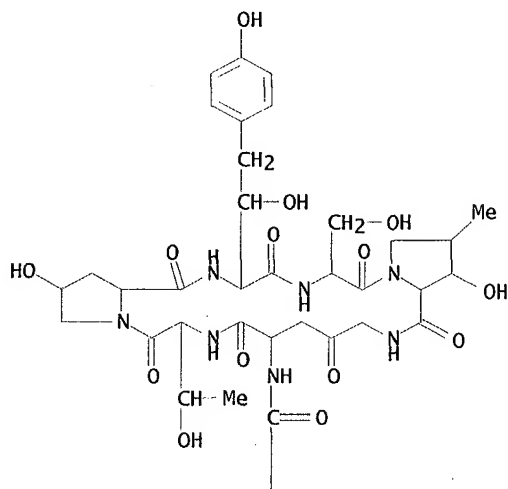




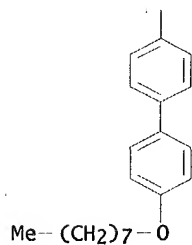
KAM 10/018,073

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PAGE 1-A



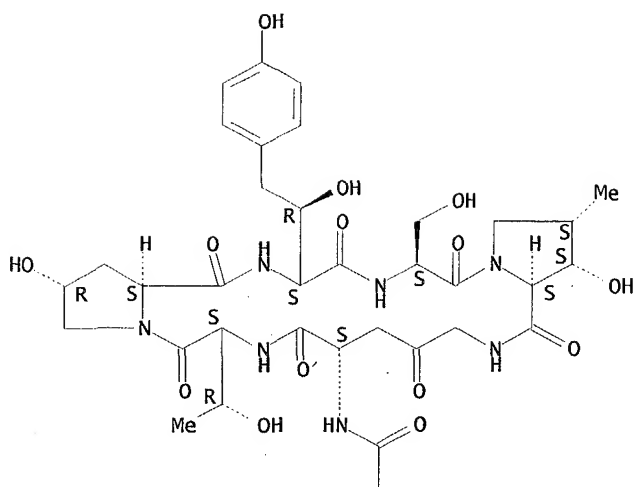
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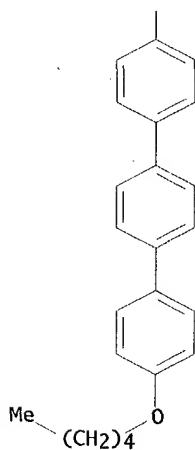
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Absolute stereochemistry.

PAGE 1-A



PAGE 2-A



IC ICM C07K007-56  
 ICS A61K038-12; A61P031-10  
 CC 34-3 (Amino Acids, Peptides, and Proteins)  
 Section cross-reference(s): 7, 10, 63  
 ST echinocandin cyclic peptide prepn antifungal glucan synthase inhibitor  
 IT Peptides, preparation  
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (cyclic; echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)  
 IT Aspergillus fumigatus  
 Fungicides  
 (echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)  
 IT 9027-19-4, Glucan synthase  
 RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL

(Biological study); PROC (Process)  
 (Candida albicans; echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)

IT 310461-86-OP 310461-89-3P 310461-95-1P  
 310461-97-3P 310461-99-5P 310462-01-2P  
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 (echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)

IT 538-75-0, N,N'-Dicyclohexylcarbodiimide 771-61-9, 2,3,4,5,6-Pentafluorophenol 5805-57-2, 2-(Aminomethyl)benzimidazole 19777-66-3  
 20439-47-8 59748-18-4 138626-63-8, Deoxymulundocandin 227472-60-8  
 RL: RCT (Reactant); RACT (Reactant or reagent)  
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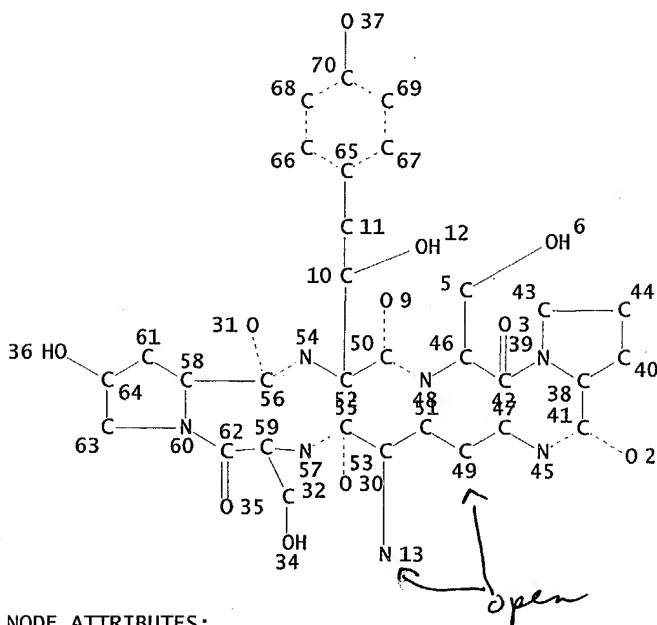
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 (echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L14

STR

parent STR



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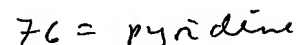
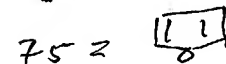
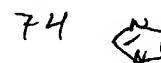
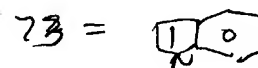
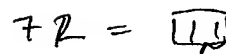
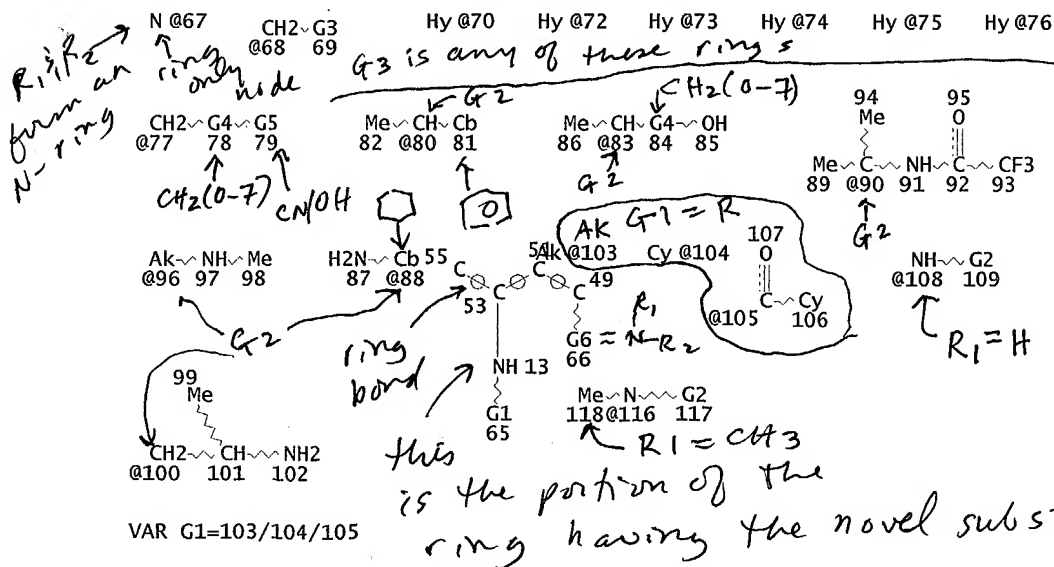
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NUMBER OF NODES IS 49

## STEREO ATTRIBUTES: NONE

L16 2586 SEA FILE=REGISTRY SSS FUL L14 2586 cpds  
L36 STR subset search



KAM 10/018,073

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RING(S) ARE ISOLATED OR EMBEDDED  
NUMBER OF NODES IS 52

STEREO ATTRIBUTES: NONE

L37 36 SEA FILE=REGISTRY SUB=L16 SSS FUL L36 36 cpds  
L38 2 SEA FILE=CAPLUS ABB=ON PLU=ON L37 2 ci + es

=> d ibib abs hitstr 138 1

L38 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2000:881187 CAPLUS

DOCUMENT NUMBER: 134:17732

TITLE: Novel echinocandin derivatives, method for preparing same and use as antifungal agents

INVENTOR(S): Corbier, Alain; Fauveau, Patrick; Pietre-Dischamp, Nathalie; Schio, Laurent; Vicat, Pascale

PATENT ASSIGNEE(S): Hoechst Marion Roussel, Fr.

SOURCE: PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: French

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
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EP 1189932	A1	20020327	EP 2000-940456	20000608

KAM 10/018,073

EP 1189932 B1 20030521  
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
IE, SI, LT, LV, FI, RO  
JP 2003504309 T2 20030204 JP 2001-502459 20000608  
AT 240971 E 20030615 AT 2000-940456 20000608  
PRIORITY APPLN. INFO.: FR 1999-7252 A 19990609  
WO 2000-FR1569 W 20000608  
OTHER SOURCE(S): CASREACT 134:17732; MARPAT 134:17732  
GI

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB The invention concerns cyclic peptides I wherein: R = chain contg. up to 30 carbon atoms, optionally contg. one or several heteroatoms, one or several heterocycles; either R1 and R2 = H, OH, alkyl optionally substituted, or NR1 forms with the carbon bearing NR1R2 a double bond and R2 is XRa, X being O, NH or N-alkyl and Ra being H, alkyl optionally substituted; R3 = H, OH, CH3; R4 = H, OH; T = H, CH3, CH2CONH2, CH2CN, (CH2)2NH2; Y = H, OH, halogen, OSO3H; W = H, OH; Z = H or CH3. The products of formula I have antifungal properties. Thus, trans-1-[4-[(2-aminocyclo-hexyl)amino]-N2-[[4-[5-[4-(pentyloxy)phenyl]-3-isoxazolyl]phenyl]carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandin B trifluoroacetate was prepd. and tested for its inhibition of glucan synthase of Candida albicans.

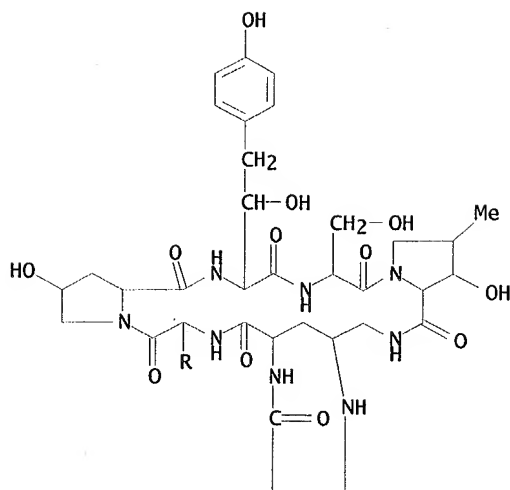
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310459-61-1P 310459-67-7P 310459-70-2P  
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(novel echinocandin derivs. method for prepg. same and use as glucan synthase inhibitors and antifungal agents)

RN 310459-20-2 CAPLUS  
CN Deoxymulundocandin, 1-[(4R)-4-[[[(1R,2R)-2-aminocyclohexyl]amino]-N2-[4-[5-[4-(pentyloxy)phenyl]-3-isoxazolyl]benzoyl]-L-ornithine]-, trifluoroacetate (salt) (9CI) (CA INDEX NAME)

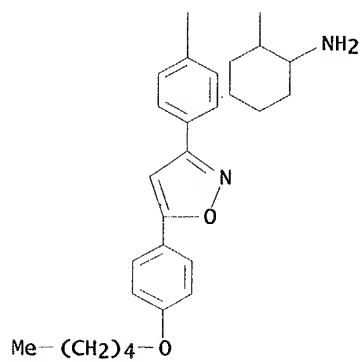
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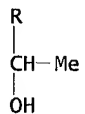
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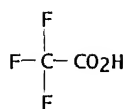
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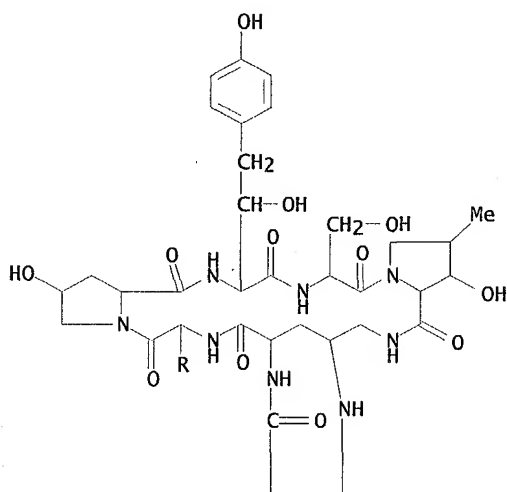


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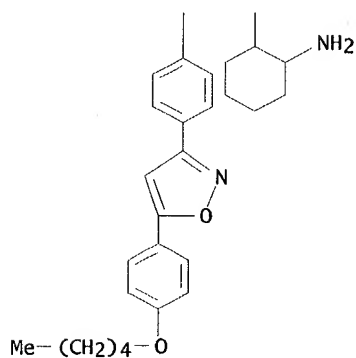
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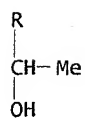
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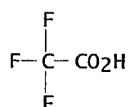






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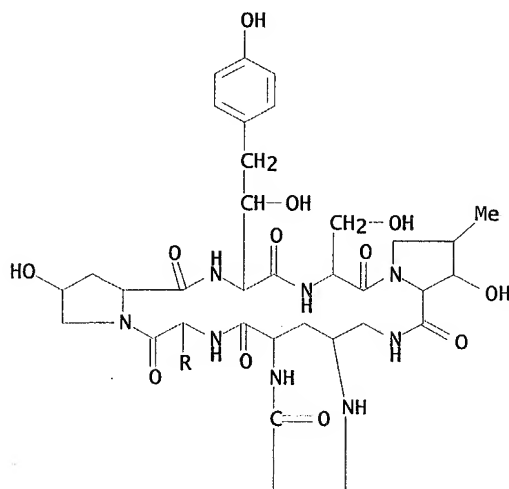
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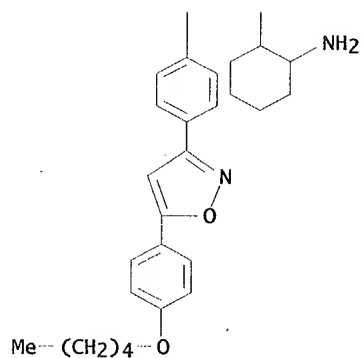
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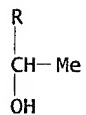


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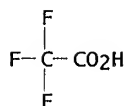


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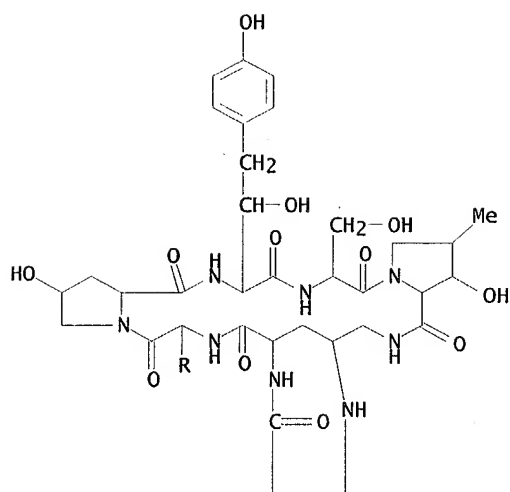


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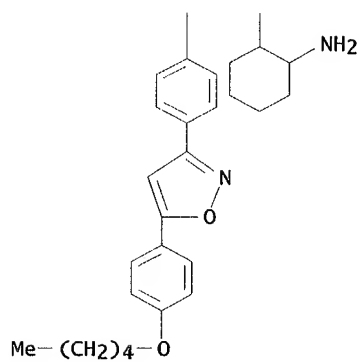
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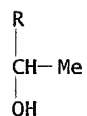
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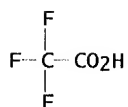
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CM 2

CRN 76-05-1  
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KAM 10/018,073



RN 310459-33-7 CAPLUS

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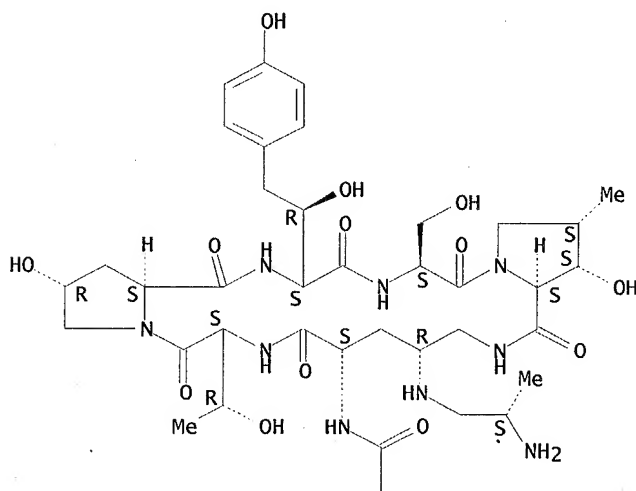
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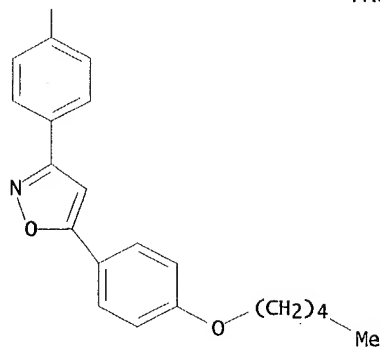
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Absolute stereochemistry.

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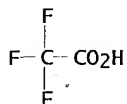
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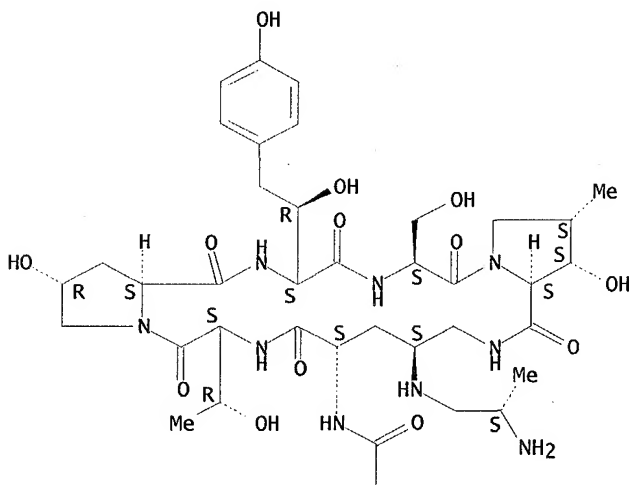
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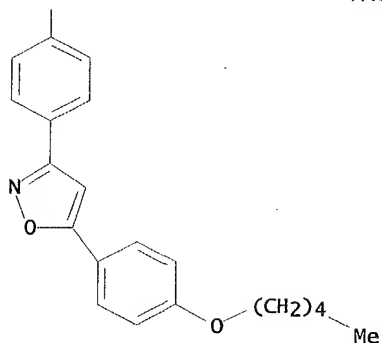
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Absolute stereochemistry.

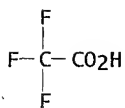
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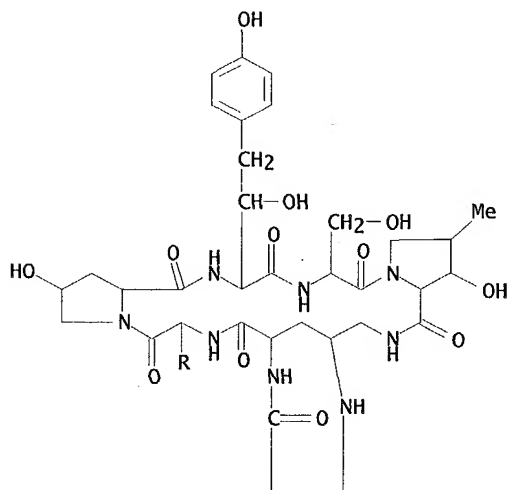
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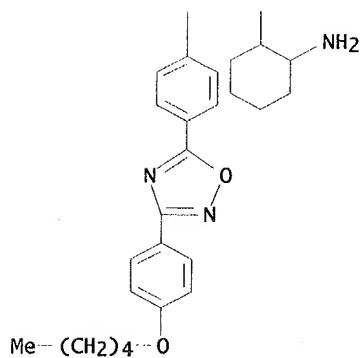
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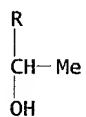


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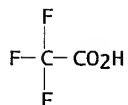


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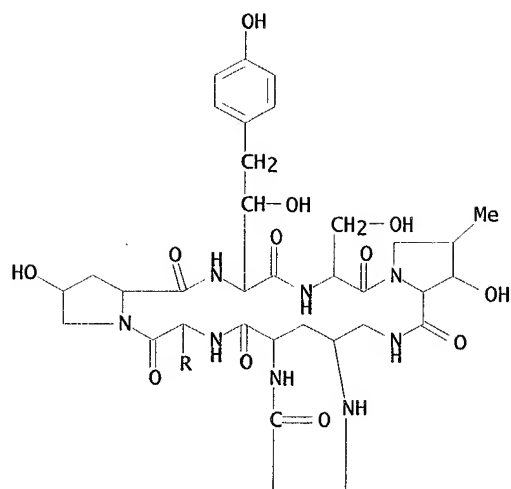


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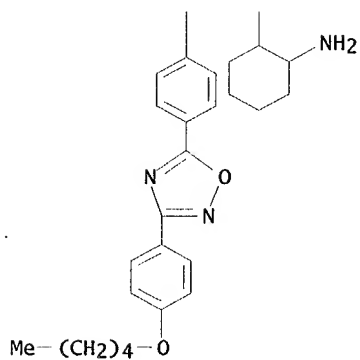
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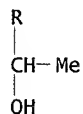
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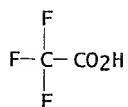


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KAM 10/018,073



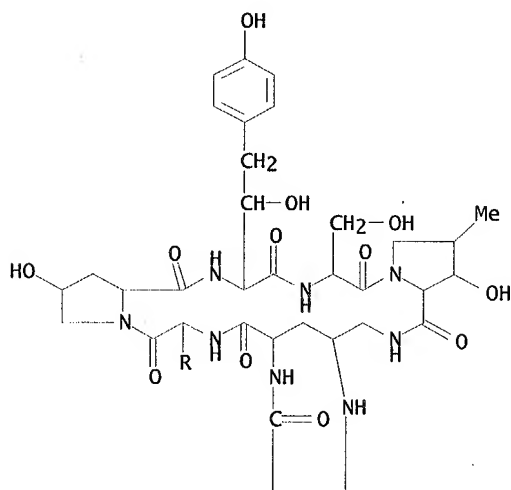
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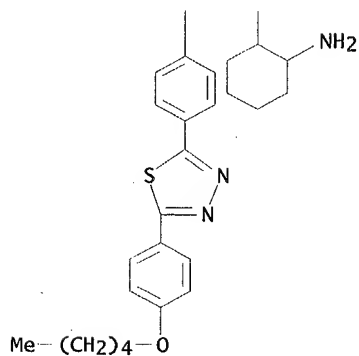
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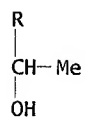
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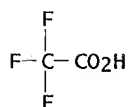
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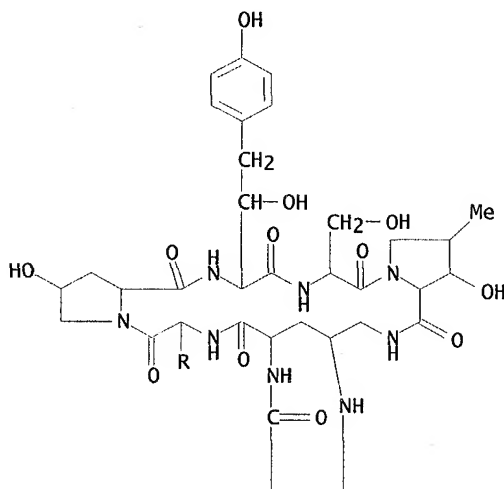
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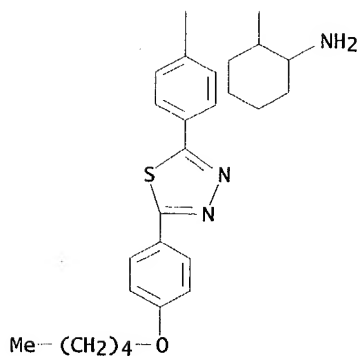
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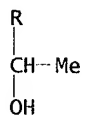


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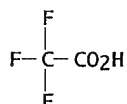


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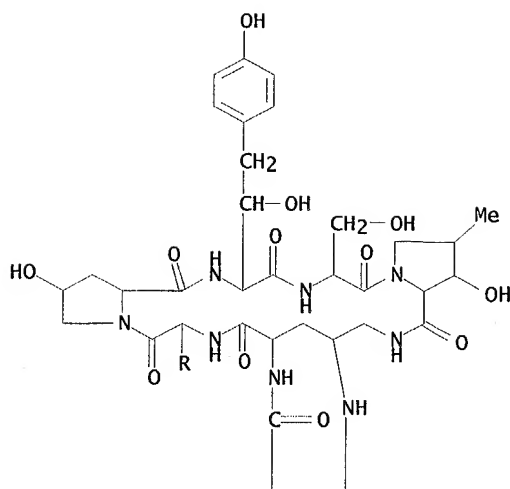


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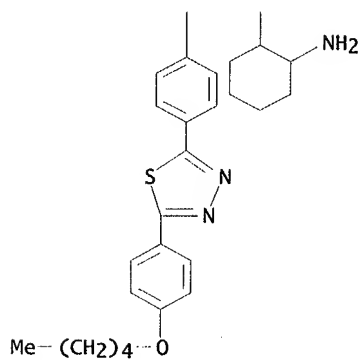
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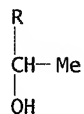
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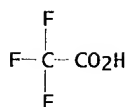
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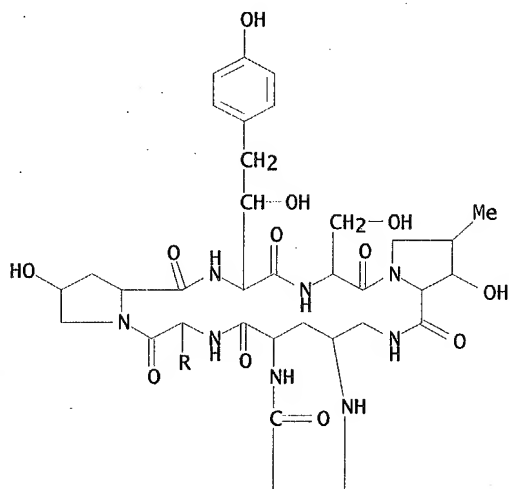
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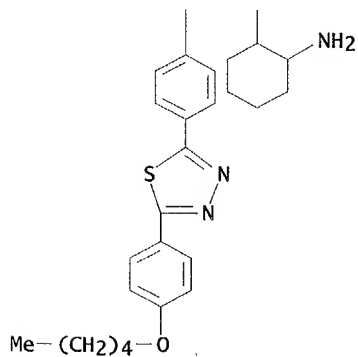
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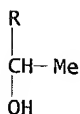
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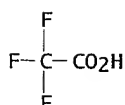


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CM 2

CRN 76-05-1  
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REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L38 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2000:881186 CAPLUS

DOCUMENT NUMBER: 134:17731

TITLE: Echinocandin derivatives, method for preparing same and application as glucan synthase inhibitors and antifungal agents

INVENTOR(S): Fauveau, Patrick; Hawser, Stephen; Lebourg, Gilles; Schio, Laurent

PATENT ASSIGNEE(S): Hoechst Marion Roussel, Fr.

SOURCE: PCT Int. Appl., 24 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: French

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

*Same citation as  
the inventor  
search*

*CPds are printed  
out to confirm  
that the search  
picked up all of  
the STR'S in the  
inventor search*

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000075177	A1	20001214	WO 2000-FR1568	20000608
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FR 2794746	B1	20021206		
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PRIORITY APPLN. INFO.: FR 1999-7251 A 19990609  
WO 2000-FR1568 W 20000608

OTHER SOURCE(S): CASREACT 134:17731; MARPAT 134:17731

GI

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB The invention concerns in all possible isomeric forms as well as their mixts., cyclic peptides I wherein: R represents a linear, branched or cyclic chain; either R1 represents H or CH3 and R2 represents cyclohexyl substituted by an amine, cyanoalkyl ; or R1 and R2 form with the nitrogen which bears them a cycle with 3, 4 or 5 carbons optionally substituted by an amine; R3 represents hydrogen, Me or hydroxyl; R4 represents hydrogen or hydroxyl; T represents hydrogen, Me, CH2CONH2, CH2CN, a (CH2)2NH2 or (CH2)2Nalk+X- radical, X being halogen and alk an alkyl radical; Y represents hydrogen, hydroxyl, halogen or OSO3H; W represents H or OH; Z represents H, CH3. The compds. of formula I have antifungal properties. Thus, . Trans 1-[4-[(2-aminocyclohexyl)amino]-N2-[4'-(pentyloxy)[1,1':4',1''terphenyl]-4-yl]carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate was prepd. and tested for its inhibition of glucan synthase of *Candida albicans* and of the enzyme prepd. from *Aspergillus fumigatus*.

IT 310461-86-0P 310461-89-3P 310461-95-1P  
310461-97-3P 310461-99-5P 310462-01-2P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
(echinocandin derivs., method for prepg. same and application as glucan synthase inhibitors and antifungal agents)

RN	310461-86-0	CAPLUS
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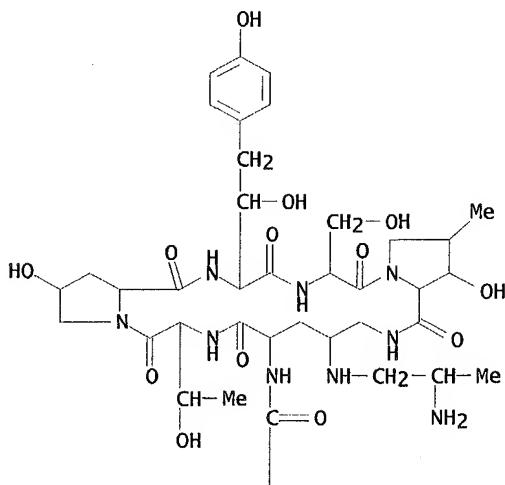
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CM 1

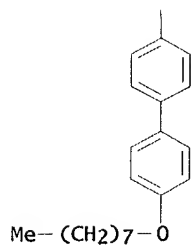
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CMF C57 H81 N9 014

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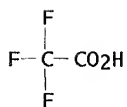


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CM 2

CRN 76-05-1  
CMF C2 H F3 O2

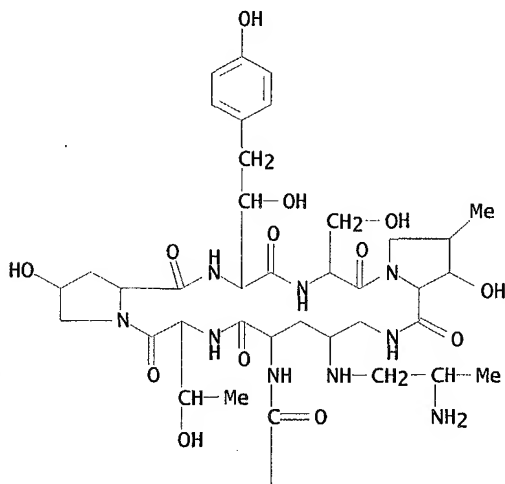


RN 310461-89-3 CAPLUS  
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CM 1

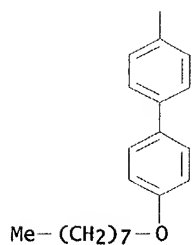
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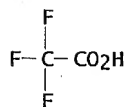


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CM 2

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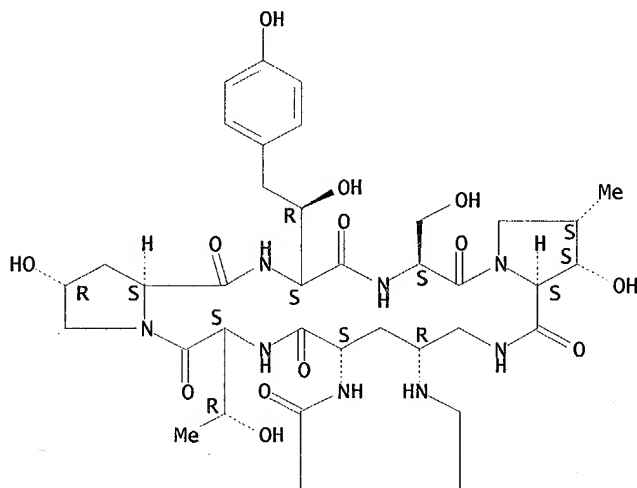
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CM 1

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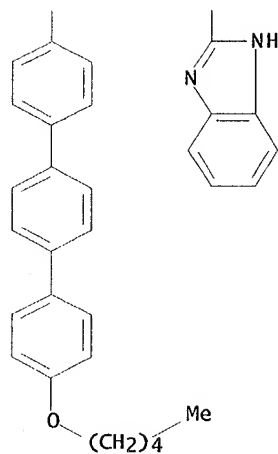
Absolute stereochemistry.

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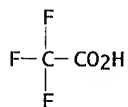
KAM 10/018,073

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CM 2

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CMF C2 H F3 O2



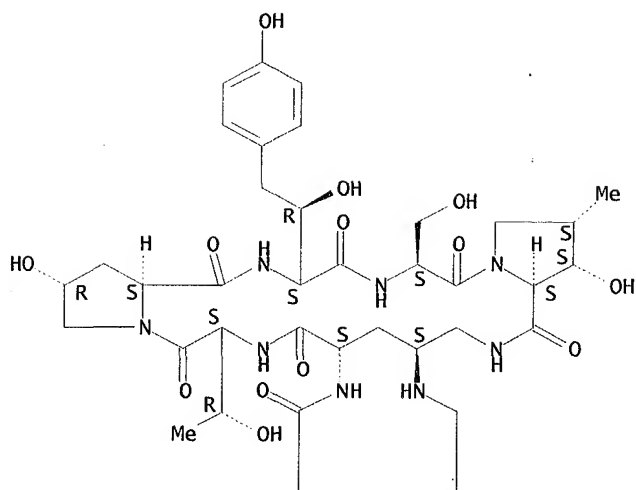
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CM 1

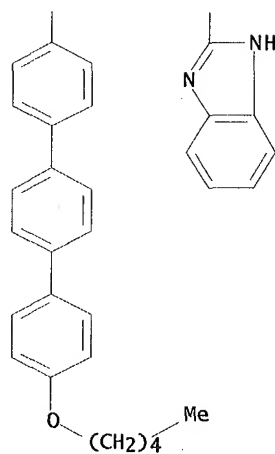
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Absolute stereochemistry.

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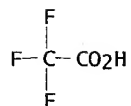


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CM 2

CRN 76-05-1  
CMF C2 H F3 O2



RN 310461-99-5 CAPLUS  
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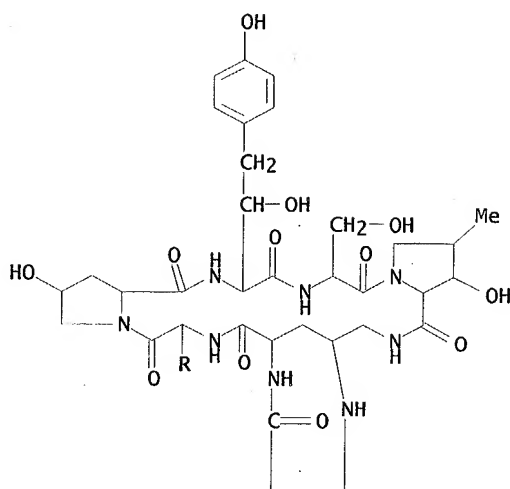
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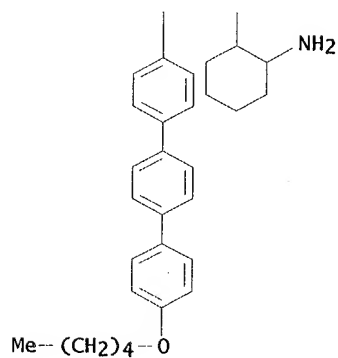
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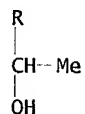
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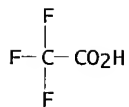
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CRN 76-05-1  
CMF C2 H F3 O2

Searched by Susan Hanley 305-4053

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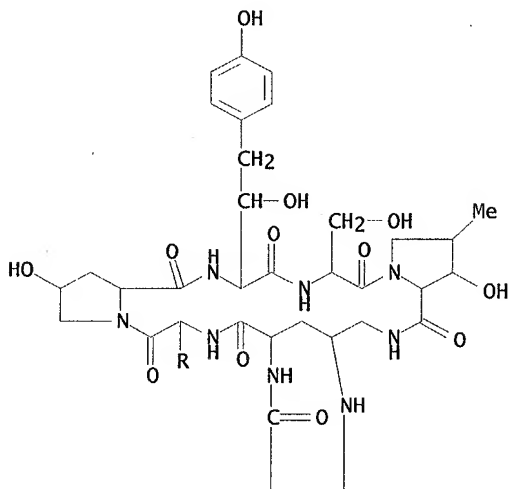


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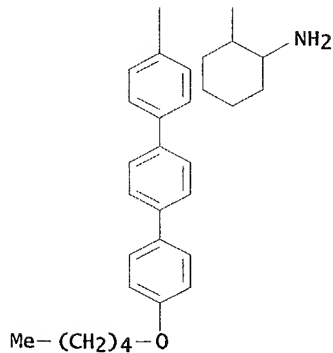
CM 1

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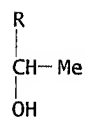


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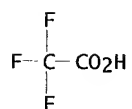
KAM 10/018,073

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CM 2

CRN 76-05-1  
CMF C2 H F3 O2



REFERENCE COUNT:

2

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